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IN THE CLAIMS:

1. (Represented original claim 1) A consumable film adapted to adhere to and dissolve in a mouth of a consumer, wherein said film comprises at least one water soluble polymer, at least one pharmaceutically active agent and at least one taste masking agent.

2. (Original) The consumable film according to claim 1, wherein said at least one water soluble polymer is a member selected from the group consisting of pullulan, hydroxypropylmethyl cellulose, hydroxyethyl cellulose, hydroxypropyl cellulose, polyvinyl pyrrolidone, carboxymethyl cellulose, polyvinyl alcohol, sodium alginate, polyethylene glycol, tragacanth gum, guar gum, acacia gum, arabic gum, polyacrylic acid, methylmethacrylate copolymer, carboxyvinyl polymer, amylose, high amylose starch, hydroxypropylated high amylose starch, dextrin, pectin, chitin, chitosan, levan, elsinan, collagen, gelatin, zein, gluten, soy protein isolate, whey protein isolate, casein and mixtures thereof.

61 3. (Original) The consumable film according to claim 2, wherein said at least one water soluble polymer is pullulan.

4. (Original) The consumable film according to claim 1, wherein said at least one pharmaceutically active agent is a member selected from the group consisting of antimicrobial agents, non-steroidal anti-inflammatory agents, antitussives, decongestants, anti-histamines, expectorants, anti-diaherals, H₂-antagonists, proton pump inhibitors, central nervous system agents, analgesics and mixtures thereof.

S/N: 09/535,005
Filed: 3/23/00
Ref. A259-03-BHJ

5. (Original) The consumable film according to claim 4, wherein the antimicrobial agent is a member selected from the group consisting of triclosan, cetyl pyridium chloride, domiphen bromide, quaternary ammonium salts, zinc compounds, sanguinarine, fluorides, alexidine, octonidine, EDTA and mixtures thereof.
6. (Original) The consumable film according to claim 4, wherein the non-steroidal anti-inflammatory agent is a member selected from the group consisting of aspirin, acetaminophen, ibuprofen, diflunisal, fenoprofen calcium, naproxen, tolmetin sodium, indomethacin, and mixtures thereof.
7. (Original) The consumable film according to claim 4, wherein the antitussive is a member selected from the group consisting of benzonatate, caramiphen edisylate, dextromethorphan, chlophedianol, diphenhydramine, salts thereof and mixtures thereof.
8. (Original) The consumable film according to claim 4, wherein the decongestant is selected from the group consisting of pseudoephedrine, phenylephrine, phenylpropanolamine, salts thereof and mixtures thereof.
9. (Original) The consumable film according to claim 4, wherein the anti-histamine is selected from the group consisting of brompheniramine maleate, chlorpheniramine maleate, carbinoxamine maleate, clemastine fumarate, dexchlorpheniramine maleate, diphenhydramine hydrochloride, diphenhydramine citrate, diphenylpyraline hydrochloride, doxylamine succinate, promethazine hydrochloride, pyrilamine maleate, tripeleminamine citrate, triprolidine hydrochloride and mixtures thereof.

S/N: 09/535,005
Filed: 3/23/00
Ref. A259-03-BHJ

10. (Original) The consumable film according to claim 4, wherein the expectorant is selected from the group consisting of guaifenesin, ipecac, potassium iodide, terpin hydrate and mixtures thereof.
11. (Original) The consumable film according to claim 4, wherein the anti-diarrheal is loperamide.
12. (Original) The consumable film according to claim 4, wherein the H₂-antagonist is selected from the group consisting of famotidine, ranitidine and mixtures thereof.
13. (Original) The consumable film according to claim 4, wherein the proton pump inhibitor is selected from the group consisting of omeprazole, lansoprazole, and mixtures thereof.
14. (Original) The consumable film according to claim 1, wherein the at least one taste masking agent is an ion exchange resin.
15. (Original) The consumable film according to claim 14, wherein the ion exchange resin is a sulfonated polymer comprising polystyrene cross-linked with divinylbenzene.
16. (Original) The consumable film according to claim 14, wherein the ion exchange resin is a sulfonated polymer comprising polystyrene cross-linked with 8% of divinylbenzene, with an ion exchange capacity of about 4.5 to 5.5 meq/g of dry resin (H⁺-form).

S/N: 09/535,005
Filed: 3/23/00
Ref. A259-03-BHJ

17. (Original) The consumable film according to claim 16, wherein the ion exchange resin has irregularly-shaped particles ranging in size from about 47 to about 149 micrometers.

18. (Original) The consumable film according to claim 16, wherein the ion exchange resin has spherical particles ranging in size from about 45 to about 150 micrometers.

19. (Original) The consumable film according to claim 14, wherein the ion exchange resin is a polymer composed of polystyrene cross-linked with 8% of divinylbenzene and functionalized with a quaternary ammonium group, and wherein an exchange capacity of said ion exchange resin is normally within a range of about 3 to about 4 meq/g of dry ion exchange resin.

20. (Original) The consumable film according to claim 1, wherein the at least one taste masking agent is magnesium trisilicate.

21. (Original) The consumable film according to claim 1, wherein said at least one water soluble polymer is pullulan, said at least one pharmaceutically active agent is dextromethorphan, and said at least one taste masking agent is a sulfonated polymer ion exchange resin comprising polystyrene cross-linked with divinylbenzene.

22. (Original) The consumable film according to claim 21, wherein said pullulan is present in an amount of about 40 to about 80 wt% of said film, said dextromethorphan is present in an amount of about 5 to about 40 wt% of said film, said sulfonated polymer ion exchange resin is present in an amount of about 5 to about 40 wt% of said film, and

S/N: 09/535,005
Filed: 3/23/00
Ref. A259-03-BHJ

a ratio of said dextromethorphan to said sulfonated polymer ion exchange resin is 1:3 to 3:1.

23. (Cancelled)

24. (Cancelled)

25. (Original) A method for preparing the consumable film of claim 1, said method comprising:

dissolving water-soluble ingredients in water to provide an aqueous solution;
mixing at least one water soluble film former and at least one stabilizing agent to provide a film-forming mixture;

combining said film-forming mixture and said aqueous solution to provide a hydrated polymer gel;

mixing oils to form an oil mixture;

adding said oil mixture to said hydrated polymer gel and mixing to provide a uniform gel;

casting the uniform gel on a substrate; and

drying the cast gel to provide said film.

26. (Original) The method of claim 25, wherein said at least one pharmaceutically active agent and said at least one taste masking agent are incorporated into said aqueous solution or into said uniform gel.

27. (Original) The method of claim 25, wherein said at least one taste masking agent is an ion exchange resin, and said at least one pharmaceutically active agent is sorbed

to said ion exchange resin without separating ion exchanged pharmaceutically active agent from unexchanged agent and counter ion salts.

28. (Previously Added) A consumable film adapted to adhere to and dissolve in a mouth of a consumer, wherein said film comprises at least one water soluble polymer, at least one pharmaceutically active agent and at least one taste masking agent, wherein said at least one taste masking agent comprises an ion exchange resin; and wherein said taste masking agent is in a weight ratio to said pharmaceutically active agent of about 2:1 to about 1:2.

29. (Previously Added) The consumable film according to claim 28, wherein said ratio is about 1:1.

30. (Previously Added) A consumable film adapted to adhere to and dissolve in a mouth of a consumer, wherein said film comprises at least one water soluble polymer, at least one pharmaceutically active agent and an ion exchange resin; wherein said ion exchange resin is in a weight ratio to said pharmaceutically active agent of about 2:1 to about 1:2.

61 31. (Previously Added) The consumable film according to claim 30, wherein said ratio is about 1:1.

32. (Represented, formerly amended claim 1) A consumable film adapted to adhere to and dissolve in a mouth of a consumer, wherein said film comprises at least one water

S/N: 09/535,005
Filed: 3/23/00
Ref. A259-03-BHJ

soluble polymer, at least one pharmaceutically active agent and at least one taste

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and* masking agent, wherein said at least one taste masking agent is selected from the group consisting of a magnesium trisilicate, acrylic copolymers, cellulose ethers, cellulosics, ethyl cellulose and combinations thereof.
